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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/569,862	06/13/2006	Per Holm	134391.00114	2548
64574 BLANK ROME	7590 10/22/201 E LLP	EXAMINER		
ONE LOGAN S	-	YOUNG, MICAH PAUL		
PHILADELPH	IA, PA 19103		ART UNIT	PAPER NUMBER
			1618	
			MAIL DATE	DELIVERY MODE
			10/22/2010	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Advisory Action Before the Filing of an Appeal Brief

Application No.	Applicant(s)	
10/569,862	HOLM ET AL.	
Examiner	Art Unit	
MICAH-PAUL YOUNG	1618	

	MICAH-PAUL YOUNG	1618	
The MAILING DATE of this communication appe	ars on the cover sheet with the	correspondence add	ress
THE REPLY FILED <u>07 September 2010</u> FAILS TO PLACE THIS		-	
1. The reply was filed after a final rejection, but prior to or on application, applicant must timely file one of the following rapplication in condition for allowance; (2) a Notice of Appe for Continued Examination (RCE) in compliance with 37 C periods:	the same day as filing a Notice of replies: (1) an amendment, affidat al (with appeal fee) in compliance	Appeal. To avoid abar vit, or other evidence, we with 37 CFR 41.31; or	which places the r (3) a Request
 a) The period for reply expires 3 months from the mailing date b) The period for reply expires on: (1) the mailing date of this Adno event, however, will the statutory period for reply expire land 	dvisory Action, or (2) the date set forth ter than SIX MONTHS from the mailin	ng date of the final rejection	on.
Examiner Note: If box 1 is checked, check either box (a) or (I MONTHS OF THE FINAL REJECTION. See MPEP 706.07(f).		
Extensions of time may be obtained under 37 CFR 1.136(a). The date of have been filed is the date for purposes of determining the period of extrunder 37 CFR 1.17(a) is calculated from: (1) the expiration date of the set forth in (b) above, if checked. Any reply received by the Office later may reduce any earned patent term adjustment. See 37 CFR 1.704(b). NOTICE OF APPEAL	ension and the corresponding amoun hortened statutory period for reply ori	of the fee. The appropria ginally set in the final Office	ate extension fee be action; or (2) as
2. The Notice of Appeal was filed on A brief in compl filing the Notice of Appeal (37 CFR 41.37(a)), or any exter Notice of Appeal has been filed, any reply must be filed with the complete of Appeal has been filed.	sion thereof (37 CFR 41.37(e)), t	o avoid dismissal of the	
<u>AMENDMENTS</u>			
3. The proposed amendment(s) filed after a final rejection, be (a) They raise new issues that would require further cor (b) They raise the issue of new matter (see NOTE below (c) They are not deemed to place the application in bett	isideration and/or search (see NC w);	TE below);	
appeal; and/or (d) They present additional claims without canceling a c			
NOTE: (See 37 CFR 1.116 and 41.33(a)).	, ,	,	
4. The amendments are not in compliance with 37 CFR 1.12	1. See attached Notice of Non-Co	ompliant Amendment (l	PTOL-324).
5. Applicant's reply has overcome the following rejection(s):	See Continuation Sheet.		
6. Newly proposed or amended claim(s) would be all non-allowable claim(s).	owable if submitted in a separate,	timely filed amendmer	nt canceling the
7. For purposes of appeal, the proposed amendment(s): a) [how the new or amended claims would be rejected is prov The status of the claim(s) is (or will be) as follows: Claim(s) allowed:		ill be entered and an e	xplanation of
Claim(s) allowed Claim(s) objected to: Claim(s) rejected: <u>1-10,20-25,27-29,31-37,40-44,51 and 5</u>	2		
Claim(s) withdrawn from consideration:	<u>z</u> .		
AFFIDAVIT OR OTHER EVIDENCE	before or on the data of filing a N	latice of Annaal will not	t be entered
 The affidavit or other evidence filed after a final action, but because applicant failed to provide a showing of good and was not earlier presented. See 37 CFR 1.116(e). 	sufficient reasons why the affida	vit or other evidence is	necessary and
 The affidavit or other evidence filed after the date of filing a entered because the affidavit or other evidence failed to or showing a good and sufficient reasons why it is necessary 	vercome <u>all</u> rejections under appe and was not earlier presented. S	al and/or appellant fail: See 37 CFR 41.33(d)(1	s to provide a).
10. ☐ The affidavit or other evidence is entered. An explanation REQUEST FOR RECONSIDERATION/OTHER	of the status of the claims after e	entry is below or attach	ed.
11. The request for reconsideration has been consideration because: See Continuation Sheet.	ered but does NOT place the appl	ication in condition for a	allowance
12. ☑ Note the attached Information <i>Disclosure Statement</i> (s). (PTO/SB/08) Paper No(s) 9/7/10	12/9/09	
13. ☑ Other: See Continuation Sheet.		0, 00	
/Michael G. Hartley/ Supervisory Patent Examiner, Art Unit 1618	/MICAH-PAUL YOUN		

Continuation of 5. Applicant's reply has overcome the following rejection(s): obviousness type double patenting of claims 1-44 and 51 over claims 59,66,72-74,83-85 and 90 of copending 10/574,125 and claims 1,3-11,13-29,31-34,36,37,40,44 and 53-56 of copening 10/569,863. Copending 11/885,992 has been abandoned so the rejection has been withdrawn.

Continuation of 11. does NOT place the application in condition for allowance because: The prior art continues to obviate the instant claims by disclosing a tablet comprising tacrolimus, polyethylene glycol and Poloxamer. The '942 patent discloses a tablet formulation comprising tacrolimus and polyethylene glycol and is silent to Poloxamer. The '939 patent discloses capsule or tablet formulations comprising water insoluble drugs (tacrolimus is water insoluble) in combination with polyethylene glycol and poloxamers. The combination of carrier compounds improves the solubility of the poorly water soluble drug and would have been an obvious combination in order to improve the solubility and bioavailability of the '942 formulation. Applicant argues that the art cannot be combined since the '942 patent discloses a fast release formulation and the '939 patent discloses a sustained release formulation. However these fast and sustained release labels are relative to the individual dissolution test given. The formulations are similar in that they both provide water insoluble drug in combination with polyethylene glycol of the same molecular weight range in tablet form. The combination would have been obvious with the '939 patent providing an improvement to the formulation. Application also argues that the artisan of ordinary skill would not have been able to form tablets using the '942 formulation. Applicant argues that the components would not be ideal for compression tableting, i.e. punch-die systems. However it is the position of the Examiner that the artisan of ordinary skill would recognize this limitation and simply use another tableting method such as molding or extrusion. The prior art nor the claim specifically recite tableting method steps beyond forming the tablet. As such any method that results in a tablet comprising tacrolimus polyethylene glycol and Poloxamer would meet the limitations of the claims. Various means of tableting are well-known in the art. For these reasons the claims remain obviated .

Continuation of 13. Other:

In view of the papers filed September 7, 2010, it has been found that this nonprovisional application, as filed, through error and without deceptive intent, improperly set forth the inventorship, and accordingly, this application has been corrected in compliance with 37 CFR 1.48(a). The inventorship of this application has been changed by deletion of the following previously unnamed inventors of this application: Tomas Norling

The application will be forwarded to the Office of Initial Patent Examination (OIPE) for issuance of a corrected filing receipt, and correction of Office records to reflect the inventorship as corrected